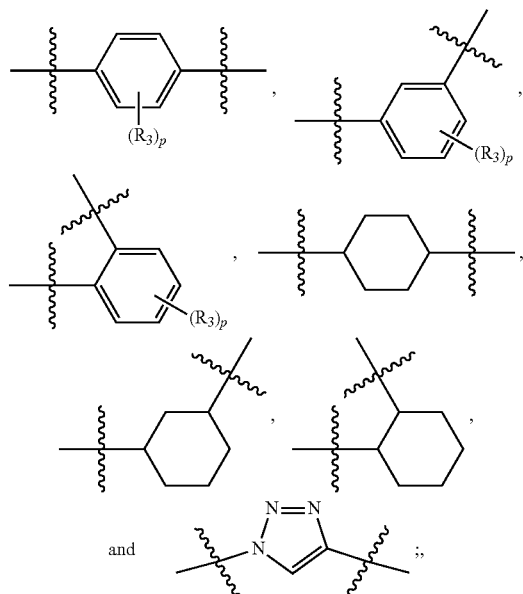


2. The compound represented by Formula I or Formula I' according to claim 1, a salt or a solvate thereof, wherein:

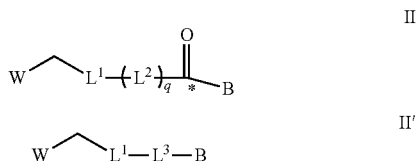
$L^1$  is selected from the group consisting of:  $-(CH_2)_m-$ ,  $-(CH_2)_tO-$ ,  $-(CH_2CH_2O)_r-$ ,  $-O-$ ,  $-NH-$ ,  $-S-$ ,  $-S(O)-$ ,  $-S(O)_2-$ ,  $-NCH_3-$ ,  $-NH(CH_2)_2NH-$ ,  $-C(O)-$ ,



wherein  $m$ ,  $t$  and  $r$  are each independently selected from the group consisting of 0, 1, 2, 3, 4, 5, 6, 7, 8, 9, 10 and 11 or

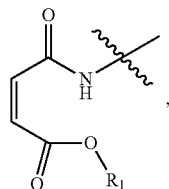
$L^1$  is  $-(CH_2)_e-C(O)NH-(CH_2CH_2O)_f-(CH_2)_g-$ , or  $-(CH_2)_h-C(O)NH-CH[(CH_2)_i-NHC(O)-(CH_2CH_2O)_j-(CH_2)_k-CH_3]-$ , wherein  $e$ ,  $f$ ,  $g$ ,  $h$ ,  $i$ ,  $j$  and  $k$  are each independently selected from the group consisting of 0, 1, 2, 3, 4, 5, 6, 7, 8, 9, 10 and 11.

3. A compound represented by Formula II or Formula II', a salt or a solvate thereof,



wherein:

W is



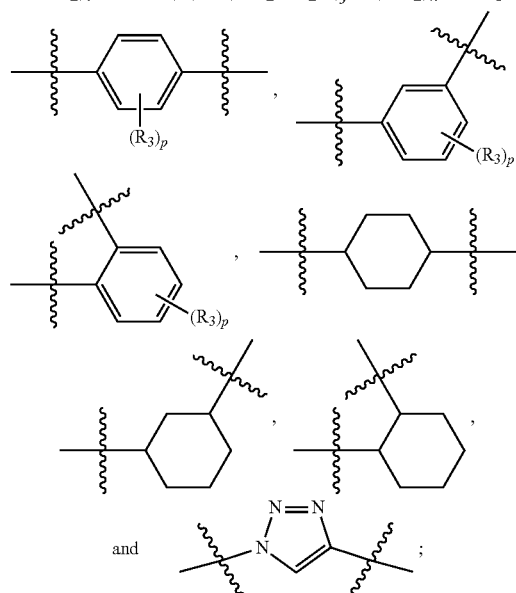
wherein the two carbonyl groups are located on the same side of the  $C=C$  double bond, which is a cis structure;

B is an active compound selected from the group consisting of drug, cytotoxin, detection reagent, diagnostic reagent and targeting carrier.

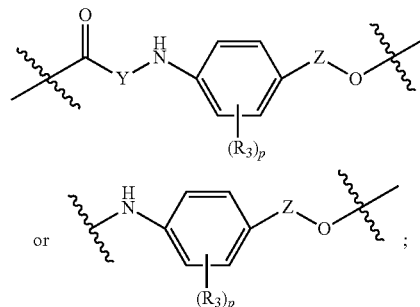
B is coupled to the site \* through a N atom or O atom in the active compound molecule; or B is coupled to  $L^3$  through a N atom or O atom in the active compound molecule;

$R_1$  is a  $C_{1-6}$  linear or branched alkyl and  $R_i$  is optionally mono- or multi-substituted by one or more substituents selected from the group consisting of: halogen and  $C_{1-4}$  alkoxy;

$L^1$  is selected from the group consisting of:  $-(CH_2)_m-$ ,  $-(CH_2)_tO-$ ,  $-(CH_2CH_2O)_r-$ ,  $-O-$ ,  $-NH-$ ,  $-S-$ ,  $-S(O)-$ ,  $-S(O)_2-$ ,  $-NCH_3-$ ,  $-NH(CH_2)_2NH-$ ,  $-C(O)-$ ,  $-(CH_2)_e-C(O)NH-(CH_2CH_2O)_f-(CH_2)_g-$ ,  $-(CH_2)_h-C(O)NH-CH[(CH_2)_i-NHC(O)-(CH_2CH_2O)_j-(CH_2)_k-CH_3]-$ ,



$L^2$  is



$L^3$  is

